

Thermal impact of magmatism in subduction zones

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Magmatism in subduction zones builds continental crust and causes most of Earth’s subaerial volcanism. The production rate and composition of magmas are controlled by the thermal structure of subduction zones. A range of geochemical and heat flow evidence has recently converged to indicate that subduction zones are hotter than predicted by models of solid-state mantle creep. We show that this discrepancy can be resolved by consideration of the heat transported by magma itself. In our one- and two-dimensional numerical models and scaling analysis, magmatic transport of sensible and latent heat locally alters the thermal structure of canonical models by ~ 300 K, increasing predicted surface heat flow and mid-lithospheric temperatures to observed values. We find the advection of sensible heat to be significantly larger than the deposition of latent heat. Based on these results we conclude that thermal transport by magma migration affects the chemistry and the location of arc volcanoes.

Petrological estimates of temperature conditions in both continental and oceanic subduction zones are systematically higher than predicted by thermal models, typically by 200–300 K [1–3]. Similarly, measurements of geothermal heat flow in SW Oregon and NE Japan are higher than predicted [1, 4] by approximately 50–100 mW/m² near the volcanic arc. Geophysical evidence from seismic imaging and magnetotelluric imaging of high temperatures and/or magma at depth under volcanic arcs [5, 6] is consistent with the emerging consensus that subduction zones are hotter than canonical models predict.

In canonical models, the thermal structure of subduction zones is calculated as a balance between diffusion of heat and advection of heat by the creeping solid mantle within the wedge-shaped region between the subducting slab and overriding lithosphere [7]. Previous modelling efforts to resolve the conflict with observations have involved varying the prescribed geometry of subduction, the coupling between mantle and slab, and the rheological model of the mantle [1, 4]. Inclusion of frictional heating along the slab top in the seismogenic zone increases heat flow in the fore-arc [8]. None of these efforts

have been successful in explaining both the amplitude of the thermal observations and their position relative to the volcanic arc.

It is known that hydrous fluids are released from the subducting slab by de-volatilization reactions [9] and percolate upward into the mantle wedge. There they reduce the solidus temperature, promote melting, and hence become silicic as they ascend. During their ascent, the fluids traverse from cooler mantle adjacent to the slab, to hotter mantle at the core of the wedge, to cooler mantle at the base of the lithosphere. They advect their temperature between these regions and consume or supply latent heat with melting and freezing. Despite the copious production of magma in subduction zones, these processes have been neglected from previous models. The circulation of hydrous fluids in the crust is believed to play a role in cooling the slab in the fore-arc region [11], and a scaling argument comparing advective heat transport by magma flow to thermal diffusion in terms of a Péclet number suggests that magma flow may also be thermally significant [10]. In this paper we assess the role of magmatic processes in altering the thermal structure of the wedge and lithosphere. Our approach is based on the theory of two-phase dynamics of the magma–mantle system [12]. We quantify the role of melts in terms of advection by melt transport and latent heat sources/sinks associated with freezing and melting reactions.

Magma migration in the mantle is a particular case of two-phase flow, governed by continuum equations of mass and momentum conservation with constitutive relations for the shear and bulk viscosity of the mantle and the mobility of the magma [12]. In order to calculate the thermal effect of melt migration, we also solve a heat equation derived from the conservation of energy, and equations for conservation of chemical species. We specify a petrological model of phase change to determine the rate at which latent heat is released by melting and freezing reactions. We omit most of these governing equations, which can be found in, for example, references [12, 13], and are discussed further in the Supplementary Material. However, since it is central to our study, we briefly describe the heat equation:

$$\frac{\partial T}{\partial t} + \bar{\mathbf{v}} \cdot \nabla T = \kappa \nabla^2 T + \frac{L}{\rho c_p} \Gamma, \quad (1a)$$

$$\bar{\mathbf{v}} = \mathbf{v}_s(1 - \phi) + \mathbf{v}_l\phi \equiv \mathbf{v}_s + \mathbf{v}_D. \quad (1b)$$

T denotes temperature, t time, κ thermal diffusivity, ρ

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density, c_p specific heat capacity, L latent heat, and Γ melting rate. We neglect differences between the thermal properties of the phases since these do not affect the solution at leading order. The velocity variables involved are: solid mantle velocity \mathbf{v}_s , liquid magma velocity \mathbf{v}_l , the Darcy (or segregation) flux $\mathbf{v}_D \equiv \phi(\mathbf{v}_l - \mathbf{v}_s)$, and phase averaged velocity $\bar{\mathbf{v}}$.

In the absence of magma, eqn. (1a) reduces to the heat equation used in canonical single-phase flow calculations. In the presence of magma, two additional terms are present. First, a latent heat sink associated with melting ($\Gamma > 0$), which becomes a source in the case of freezing ($\Gamma < 0$). Second, an additional advective term associated with the segregation flux of magma (caused, for example, by its tendency to rise buoyantly). For the latter, the decomposition of the phase-averaged velocity $\bar{\mathbf{v}}$ into \mathbf{v}_s and \mathbf{v}_D clarifies the advective role of magmatism in the energy balance of subduction zones. Hence we emphasize that the latent heat of phase change is not the only contribution; we next consider the importance of latent heat relative to advective transport by the magma.

So-called ‘melting-column models’ have been used to understand mid-ocean ridge magmatism, where the main cause of melting is decompression of the upwelling magma and mantle [14–16]. Subduction zones are a considerably more complex environment, but we adapt ideas from melting-column models to investigate how magmatism modifies subduction thermal structure. One important difference from mid-ocean ridges is that hydrous flux-melting is the dominant cause of melting in subduction zones. Therefore, we extend the two-component phase diagrams used previously [16] by adding a third, hydrous component [17]. We use a quasi-linear solidus depression with water content and choose parameters to mimic the main features of the isobaric productivity curves given by [18], which are based on compiled laboratory data. Further details of our parameterization are given in the Supplementary Material.

A one-dimensional, steady-state heat equation, with lengths rescaled by the height of the column H and velocities rescaled by the diffusive scale κ/H , can be written

$$\text{Pe } T' - \Psi = T'' - \text{Pe St } (T' + \Delta T_H), \quad (2)$$

where a prime denotes a derivative with respect to height (e.g., T' is a rescaled vertical heat flux) and ΔT_H is melting rate associated with decompression. Two dimensionless numbers control the behaviour of the system: a Péclet number $\text{Pe} = H\bar{W}_0/\kappa$ is the scaled volume flux at the base of the column and quantifies the flux of fluid off the subducting slab; and a Stefan number $\text{St} = (L/c_p)\partial F/\partial T$ is a scaled isobaric productivity and quantifies the latent heat of melting.

The mantle flow in subduction zones is far from one-dimensional; a corner flow is driven by the motion of the subducting slab [7]. A key step in representing corner flow in a column model is to introduce a spatially variable volumetric heating term Ψ that mimics the effects of large-scale mantle advection. We calculate Ψ from

a single-phase, two-dimensional flow and thermal structure model, using the domain geometry and temperature-dependent viscosity given by a benchmark study [19]. We show the geometry and thermal structure in the Supplementary Material. This is generic, not intended to represent any particular subduction zone. We extract a vertical temperature profile at some position of interest $T_{\text{ref.}}(z)$, and use it to calculate the source term $\Psi = -T''_{\text{ref.}}$. In the absence of fluid flow ($\text{Pe} = 0$), this model returns $T = T_{\text{ref.}}$. For $\text{Pe} > 0$, this approach is reasonable provided melt does not drastically change the large-scale mantle dynamics, a prospect we consider below.

We present example column solutions in Fig. 1. These use a slab-fluid source at 100 km depth, roughly consistent with the observed mean position of arc volcanoes [21, 22]. The hydrous flux at the base of the column is varied within the range suggested by a previous study of fluid flow in subduction zones [20]. Column (a) shows profiles of the absolute temperature; column (b) shows the temperature difference compared to a single-phase (magma-free) reference case. The change in temperature from the reference state increases with fluid transport, and is significant even at the lower end of the range of fluxes [20]. Immediately above the slab, fluid flow reduces the mantle temperature as material is transported from the relatively cold slab. Nearer the surface, the effect is reversed as fluid flow brings warm material from the mantle into the lithosphere. This effect is supplemented by latent heat release, shown in column (c). Above the slab, melting of the mantle wedge facilitated by the presence of water consumes latent heat. Nearer the surface, solidification of the melt deposits latent heat. It is interesting to note that the maximum degree of melting (d) does not vary monotonically with fluid flow, but peaks at an intermediate Péclet number between 2 and 5.

The main physical mechanism giving rise to this temperature signal is advection by the magma, represented in the advection term of eqn. (2). Indeed, the latent heat of freezing/melting is relatively small. In the Supplementary Material, we report further calculations that show that neglecting latent heat makes little difference to the temperature profiles, and that changing the parameterization of hydrous flux melting (either to mimic more closely a more detailed parameterization [18], or by artificially doubling the Stefan number) is also inconsequential.

Two-dimensional effects that are neglected in column models such as lateral diffusion, changes to mantle flow, and sub-vertical magma flow require a more careful treatment. Our approach is to modify the reference simulation of the thermal structure of a subduction zone [19] by applying a prescribed segregation flux \mathbf{v}_D in the heat eqn. (1a). We assume that magma segregates purely vertically, driven by the density difference between solid and liquid phases; and prescribe Gaussian fluid flow profiles centred at the typical position of the arc volcano [21, 22]. Our numerical scheme solves iteratively for ther-

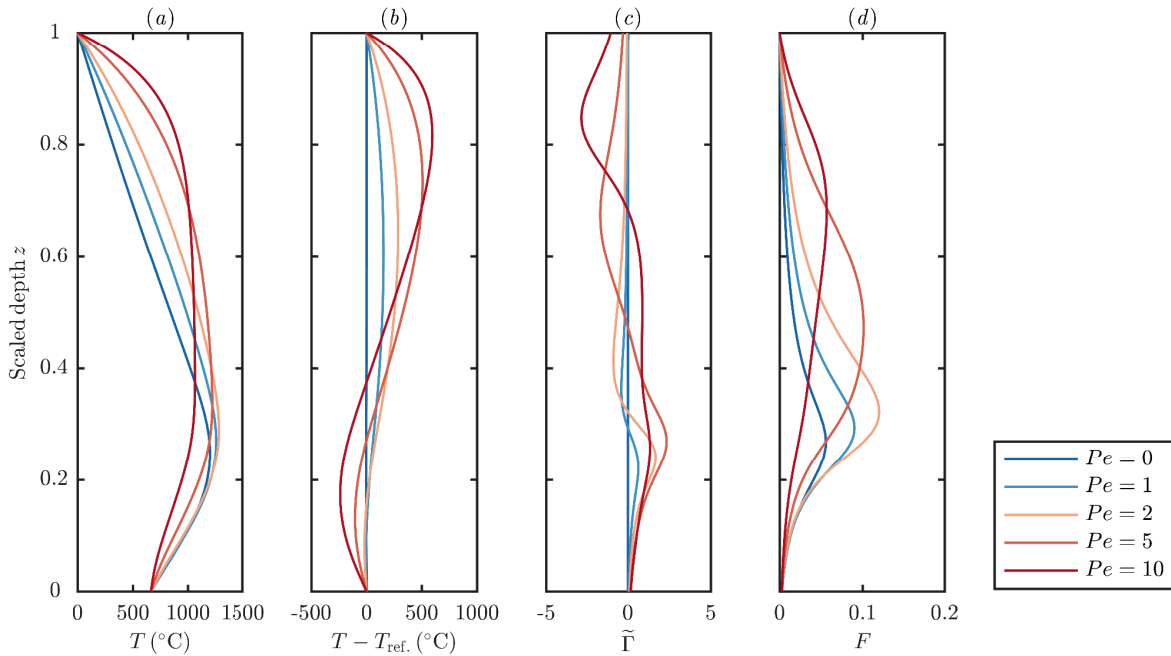


FIG. 1. Melting column model with prescribed temperature at the base ($z = 0$) and top ($z = 1$) of the melting column, which runs from the slab to the surface respectively. (a) temperature profiles, (b) temperature perturbation caused by magmatism, (c) scaled melting rate, (d) degree of melting. The range of Péclet number considered is roughly equivalent to melt fluxes from the slab reported in Wilson *et al.* [20]. Bulk water content is 0.5%.

mal structure and solid flow, which are coupled through the temperature-dependence of mantle viscosity, until a steady-state is achieved. The thermal impact of magmatism is then defined as the difference between the calculated and reference temperature fields. Note that we do not consider any latent heat effects in this exercise because we previously observed that these are negligible.

The two-dimensional calculations predict that fluid flow substantially alters the thermal structure in subduction zones, as shown in Fig. 2. The main effect is to raise temperatures near the base of the lithosphere, where warm material is transported from the mantle upward. These 2D results are qualitatively similar to the 1D column models (cooling above the slab, warming near the surface), indicating that the physical mechanisms discussed in the previous section remain pertinent. Some features only occur in two dimensions, such as the along-slab cooling observed deeper than the fluid source that is caused by advection by the mantle flow. Thus the thermal impact of magmatism is felt beyond where the magma itself flows.

Using a reference estimate of global arc magma production of $1 \text{ km}^3/\text{yr}$ [26], comparable to the estimate used in England and Katz [27], magmatism raises temperatures by up to 270 K (Fig. 2b). We also consider a magmatic flux 50% smaller or larger than this reference case. Temperatures are raised by $\sim 150 \text{ K}$ (Fig. 2a) with the lower estimate (consistent global average estimate of arc vol-

canism excluding plutonism [28]). The higher estimate raises temperatures by up to 380 K (Fig. 2c). In the Supplementary Material, we separately consider fluid release associated with the major dehydration reactions of the slab. We also show that the coupling of the solid flow to the heat equation increases the thermal perturbation (since the solid viscosity is reduced leading to increased solid velocities, a positive feedback), that only the total flux (not the width) of the fluid source is thermally significant, and that a thicker overriding plate reduces the thermal perturbation.

Elevated near-surface temperatures in subduction zones are associated with elevated heat flow, as shown in Fig. 3. This elevated heat flow is strongest at the position of the arc, over a width of around 50 km. The width is determined by thermal diffusion rather than the imposed width of fluid source. Calculations with a magma flux between the reference and high values are consistent with heat-flow observations.

Evidence from petrology and heat flow measurements suggests that temperatures in subduction zones are some 200–300 K hotter than would be expected on the basis of mantle flow alone [1–3]. This discrepancy is observed to peak at around 60 km depth, comparable to depth where we find magmatism to have the greatest thermal impact. Fig. 4 shows that including melt migration in thermal models can reconcile much of this discrepancy. This combination of observation and thermal modelling

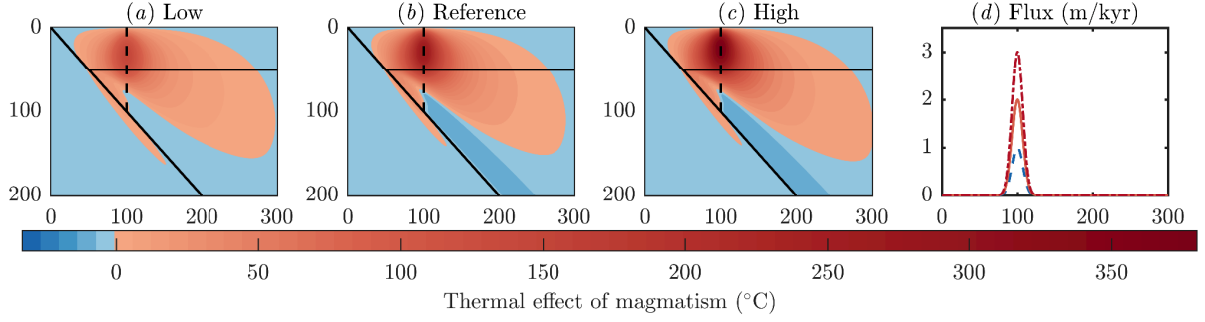


FIG. 2. The thermal impact of magmatism ($T - T_{\text{ref.}}$) associated with a source beneath the volcanic arc (dashed black line). The slab and overriding plate geometry are shown by solid black lines. We compare a low, medium and high estimate of the magmatic flux (*a-c*). The prescribed magma flow (segregation flux) is shown in (*d*). Horizontal and vertical scales are distance from the trench, in kilometres.

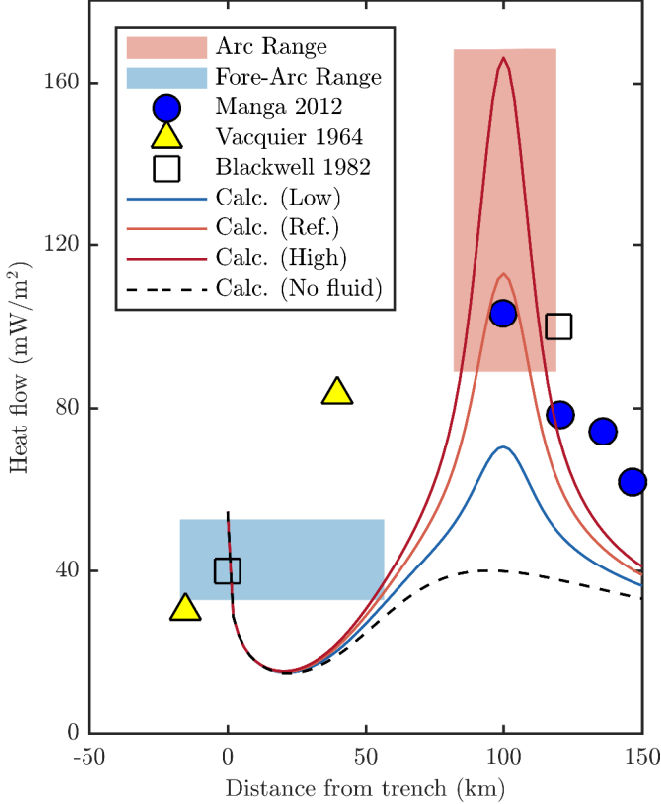


FIG. 3. Elevated heat flow in subduction zones associated with melt migration is consistent with observed ranges [23–25] in oceanic and continental subduction zones. The heat flow is raised by around 40–120 mW/m² concentrated near the region of peak fluid flow 100 km from the trench. The results were obtained by evaluating surface temperature gradients from Fig. 2 and converting to heat flow using a constant thermal conductivity of 2.52 W/m/K.

based on two-phase flow supports the hypothesis that magmatism significantly alters the thermal structure of subduction zones.

Scaling arguments also support this argument. Indeed,

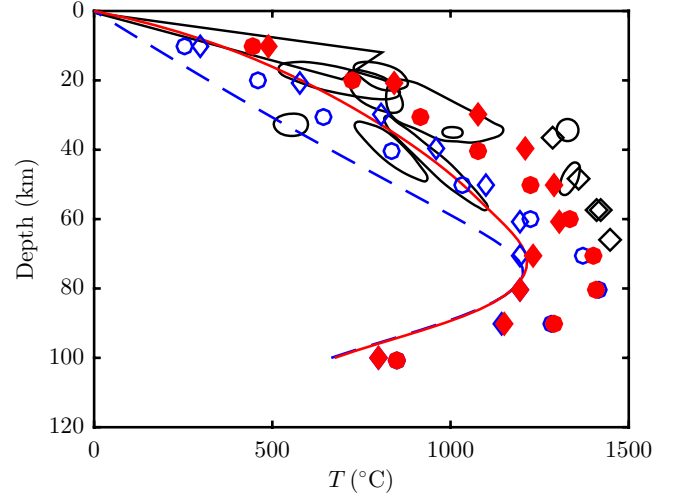


FIG. 4. Temperature structure compared to a compilation of petrological and heat flow data (black open shapes are taken from Plate 1 in Reference [1]). We shift the output of two thermal models [4, 29] by the thermal impact of melt migration calculated as the reference case in Fig. 2(*b*). The original temperatures are open blue circles and diamonds and the modified temperatures are shown in solid red markers of the corresponding shape. Our reference (blue dashed) and modified (solid red) thermal profiles are also included, but note that these were computed with a greater subduction angle from the geometry of [19].

it is possible to approximate the thermal effect of magmatism due to advection as follows. The elevated heat flow

$$Q \approx \frac{F_V \rho c_p \Delta T}{A_{\text{sub}}} \approx 80 \text{ mW/m}^2, \quad (3)$$

based on a global magma flux $F_V = 1 \text{ km}^3 \text{ yr}^{-1}$ [26], density $\rho = 3 \times 10^3 \text{ kg m}^{-3}$, heat capacity $c_p = 1.2 \times 10^3 \text{ J kg}^{-1} \text{ K}^{-1}$, $\Delta T \approx 1350 \text{ K}$, and an area of elevated heat flow $A \sim 2 \times 10^{12} \text{ m}^2$ (the total length of $50 \times 10^3 \text{ km}$ and an assumed width of 40 km). This

is consistent with Fig. 3. We can also support our contention that advective heat transport matters more than latent heat release by estimating the ratio R of these terms in eqn. (1a):

$$R \approx \frac{c_p \Delta T}{L} \frac{\rho |\mathbf{v}_D|}{\Gamma H} \approx \frac{c_p \Delta T}{L} \approx 3.2, \quad (4)$$

where $L = 5 \times 10^5 \text{ J kg}^{-1}$. We have used the fact that, at steady state, $\rho |\mathbf{v}_D| / \Gamma H \approx 1$ on average, since there is a balance between melt production, melt extraction, and melt solidification. Therefore, magmatism has a significant thermal effect and this effect is mainly due to advection by the magma. This latter finding is in contrast to a one-dimensional model of thermal erosion that balances latent heat release with vertical diffusion, but neglects advection and horizontal diffusion [2, 27].

The thermal signature of melt migration should be considered when interpreting heat flow, petrologic, gravity, and seismic data. A perturbation as large as 300 K is likely to affect the chemistry of arc volcanoes [30]. It also significantly affects the solid mantle flow through reduction of mantle viscosity, leading to increased circulation in the mantle wedge [29]. Viscosity also affects magma pathways in subduction zones through its effect on the compacting length, focussing magmas from a broader area to beneath the arc volcanoes [20, 31]. Thus, consistent with evidence from global systematics [27], coupled mantle–magma flow may well affect the location of arc volcanoes themselves.

Author contributions. D.R.J. and M.T. developed

the one-dimensional melting column model and petrological model of melting. D.R.J., R.F.K. and J.F.R. developed the two-dimensional thermal model with magmatism. J.F.R. contributed a single-phase numerical code to compute the thermal structure of a subduction zone, to which D.R.J. added two-phase flow. R.F.K. and D.R.J. compared the model with petrological and heat flow observations. D.R.J. wrote the manuscript with R.F.K., and discussed the manuscript with M.T. and J.F.R. All authors jointly discussed and analyzed the data, results, conclusions, and implications.

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SUPPLEMENTARY MATERIAL

I. PETROLOGICAL MODEL OF HYDROUS FLUX MELTING

In our melting column model, we use a simple petrological model of hydrous flux melting, which is the dominant form of melting in subduction zones. The model is developed as follows. First, we restrict attention to a ternary system. The three components should not be thought of as identifiable minerals but rather as idealized components chosen to capture the physics in which we are interested. We start with two components that can be considered ‘refractory’ and ‘fertile’ [14, 16]. To this system, we add a third component to represent volatiles. We initially will take this component to be ‘water.’ It must be understood that our model will only be valid in the case that the concentration of the ‘water’ component is relatively small. The main role of this third hydrous component is to depress the solidus temperature.

Our second simplification is to use a linear phase diagram. This can be thought of as a linearization of the ternary phase loops used by Keller and Katz [17] about some initial composition at the bottom of the melting column.

Our third simplification is that the melting/freezing reactions happen sufficiently rapidly that a partially molten region is at thermodynamic equilibrium. This implies that compositions of the solid and liquid phases are controlled directly by the phase diagram.

A. Mathematical description of phase diagram

The solidus temperature increases with increasing pressure at a rate γ . We linearize the dependence of the solidus on chemical composition. Since the sum of the concentrations of the components is unity, we need only specify 2 linear coefficients M_2, M_3 for the fertile and water components respectively, both of which lower the solidus temperature. Thus the solidus temperature

$$T_s = T_{s0} - \rho g z / \gamma - M_2 c_2^s - M_3 c_3^s. \quad (5)$$

This expression can be rearranged to give, for example, the solidus concentration c_2^s as a function of temperature, depth, and concentration of the third component. An interpretation of equation (5) can be made by identifying $T_{s0} - \rho g z / \gamma$ with the solidus temperature of the refractory component at given depth z , which is this section is negative.

We assume that the liquidus concentration is related to the solidus concentration as follows

$$c_2^l = c_2^s + \Delta c_2, \quad (6)$$

$$c_3^l = c_3^s + \Delta c_3. \quad (7)$$

For the simplest case we take $\Delta c_{2,3}$ to be constants, but we will also consider generalizations.

B. Choice of parameter values and implications for melting

We choose parameters in our model to constrain the degree of so-called ‘batch melting’ as a function of temperature and pressure:

$$F = \frac{\widetilde{c}_{0j} - c_j^s}{c_j^l - c_j^s}. \quad (8)$$

Batch melting refers to the degree of melting experienced by a sample raised to given temperature and pressure conditions assuming no extraction of melt from the batch or sample. The composition \widetilde{c}_{0j} is the composition of the solid mantle before the onset of melting. We then combine equations (5)–(7), which apply for each j , with equation (8) to obtain

$$F = \frac{\widetilde{c}_{03} - c_3^s}{\Delta c_3} = \frac{T - T_{s0} + \rho g z / \gamma + M_2 \widetilde{c}_{02} + M_3 \widetilde{c}_{03}}{M_2 \Delta c_2 + M_3 \Delta c_3}. \quad (9)$$

A key quantity is the isobaric productivity $\partial F / \partial T$. If Δc_2 and Δc_3 are constants, then the isobaric productivity is a constant

$$\frac{\partial F}{\partial T} = \frac{1}{M_2 \Delta c_2 + M_3 \Delta c_3}. \quad (10)$$

Thus melt is produced at a constant rate with increasing temperature. Linear models of two component melting already include this effect [e.g. 16].

In this formulation, volatiles do indeed depress the solidus temperature. However, as well as depressing the solidus, volatiles are also associated with a ‘low-productivity tail.’ The initial melting above the solidus temperature is less productive than later melting:

$$\left. \frac{\partial F}{\partial T} \right|_{F=0} < \left. \frac{\partial F}{\partial T} \right|_{F=1}. \quad (11)$$

The purely linear model does not satisfy this constraint, because the productivity is constant. Therefore, we generalize our model to allow for a low-productivity tail. Volatiles are incompatible, and partition into the melt with a partitioning coefficient D defined by

$$c_3^s = D c_3^l \Rightarrow \Delta c_3 = c_3^s (1/D - 1), \quad (12)$$

where $D \ll 1$ for volatile elements. We assume that D is constant. However, Δc_3 is no longer constant, instead depending on composition, and hence pressure and temperature. Upon a little rearrangement, we find

$$\begin{aligned} T - (T_{s0} - \rho g z / \gamma - M_2 \widetilde{c}_{02} - M_3 \widetilde{c}_{03}) \\ = (M_2 \Delta c_2 / \Delta c_3 + M_3) (\widetilde{c}_{03} - c_3^s), \end{aligned} \quad (13)$$

which can be rearranged to give a quadratic equation for c_3^s , recalling that Δc_3 is proportional to c_3^s . The degree of melting F is no longer a linear function (however it can be computed explicitly using the quadratic formula

so there is no computational difficulty, unlike more complex nonlinearities where iterative methods are required to solve for F). We can calculate the isobaric productivity at $F = 0$ and $F = 1$ and find

$$\left. \frac{\partial F}{\partial T} \right|_{F=0} = [M_3 \widetilde{c}_{03}(1/D - 1) + M_2 \Delta C_2]^{-1}, \quad (14)$$

$$\left. \frac{\partial F}{\partial T} \right|_{F=1} = [M_3 \widetilde{c}_{03} D(1 - D) + M_2 \Delta C_2]^{-1}. \quad (15)$$

We can interpret the effective isobaric productivity of the mixture as the harmonic mean of productivities associated with the fertile and volatile component. Typically, the contribution of the volatile component dominates at small F and the fertile component dominates at large F because

$$M_3 \widetilde{c}_{03} D(1 - D) \ll M_2 \Delta C_2 \ll M_3 \widetilde{c}_{03}(1/D - 1). \quad (16)$$

Our formulation thus achieves the low-productivity tail expected physically and recovers the fertile-refractory system in the absence of volatiles.

To summarize, the degree of melting increases over the temperature range

$$T(F = 0) = T_{s0} - \rho g z / \gamma - M_2 \widetilde{c}_{02} - M_3 \widetilde{c}_{03}, \quad (17)$$

$$T(F = 1) = T_{s0} - \rho g z / \gamma - M_2 \widetilde{c}_{02} + M_2 \Delta c_2 - M_3 D \widetilde{c}_{03}. \quad (18)$$

The gradient of the function $F(T)$ at these endpoints is given by equations (14), (15).

Thus a quite limited number of parameters can describe a significant range of realistic melting behaviours, as shown in Fig. 5. For the anhydrous fertile-refractory part of the system, we use $T_{s0} = 1100$ °C, $\rho g / \gamma = 4.5 \times 10^{-3}$ °C/m, $M_2 = 700$ °C, $\Delta c_2 = 0.6$, and $\widetilde{c}_{02} = 0.15$. This ensures that we match the anhydrous melting curve of Katz, Spiegelman, and Langmuir [18], particularly around 3 GPa. For the hydrous part of the system, we use $M_3 = 2 \times 10^5$ °C, $D = 0.01$. This choice of parameters was chosen to roughly match the hydrous melting curve of Katz, Spiegelman, and Langmuir [18], particularly around 3 GPa with 0.5 wt% water. Although the precise parameter values are in the right region for consistency with previous studies and their associated experimental libraries, as well as inferences from field observations, this parameterization is too simple to reproduce all the features observed experimentally. However, it can reproduce the two main features: solidus depression and a low-productivity tail.

C. Generalized model: accounting for saturation in water

The addition of more water does not indefinitely lower the solidus, because eventually water becomes saturated in the liquid phase. The amount of water that dissolves

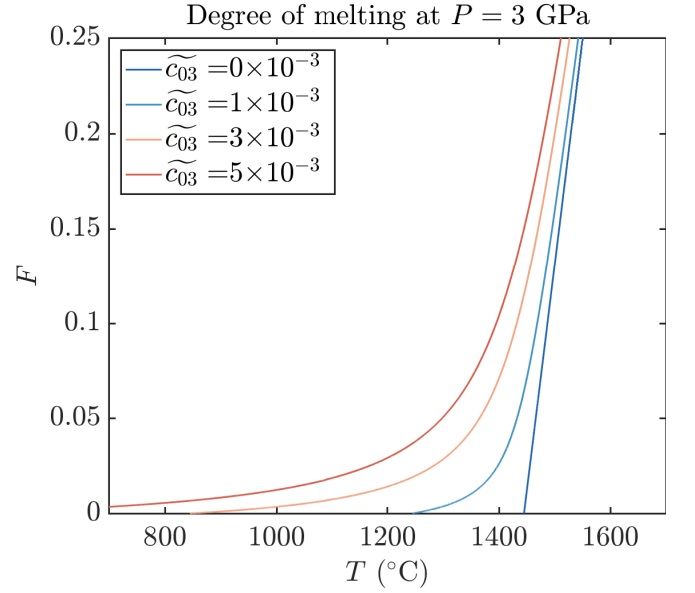


FIG. 5. The degree of melting F as a function of temperature T at increasing water concentration. Other parameters were fixed, namely $T_{s0} - \rho g z / \gamma = 1550$ °C at $z = 100$ km, $M_2 = 700$ °C, $M_3 = 2 \times 10^5$ °C, $D = 0.01$, $\Delta c_2 = 0.6$, and $\widetilde{c}_{02} = 0.15$. These numbers are motivated by Katz, Spiegelman, and Langmuir [18].

increases with pressure, and Katz, Spiegelman, and Langmuir [18] use the formula

$$X_{\text{H}_2\text{O}}^{\text{sat}} = 12.00 P^{0.6} + 1.00 P, \quad (19)$$

where the pressure P is measured in GPa. This is well constrained by experiment below 2 GPa, and constrained indirectly at higher pressures. This corresponds to a critical degree of melting and critical temperature below which the degree of melting drops rapidly to zero, as shown in Fig. 6c, for example.

Our modelling approach is to mimic this behaviour by modifying the phase diagram. We first compute the corresponding critical solid saturation point c_{sat}^s , using equation (19) for the liquid saturation, and the partition coefficient of equation (12). For temperatures below this point, we change the freezing point depression coefficient:

$$T_s = T_{s0} - \rho g z / \gamma - M_2 c_2^s - M_4 (c_3^s - c_{\text{sat}}^s) - M_3 c_{\text{sat}}^s, \quad (20)$$

where $M_4 \leq M_3$. Note that the previous model is a special case $M_4 = M_3$, and a eutectic-like phase diagram can be obtained by the special case $M_4 = 0$. In practice, we find $M_4 = M_3/50$ makes a decent approximation to Katz, Spiegelman, and Langmuir [18], as shown in Fig. 6. This means that the initial productivity near $F = 0$ is a factor $M_3/M_4 = 50$ times greater. We will use this generalized model to assess the significance of the increased productivity at volatile saturation in the following section.

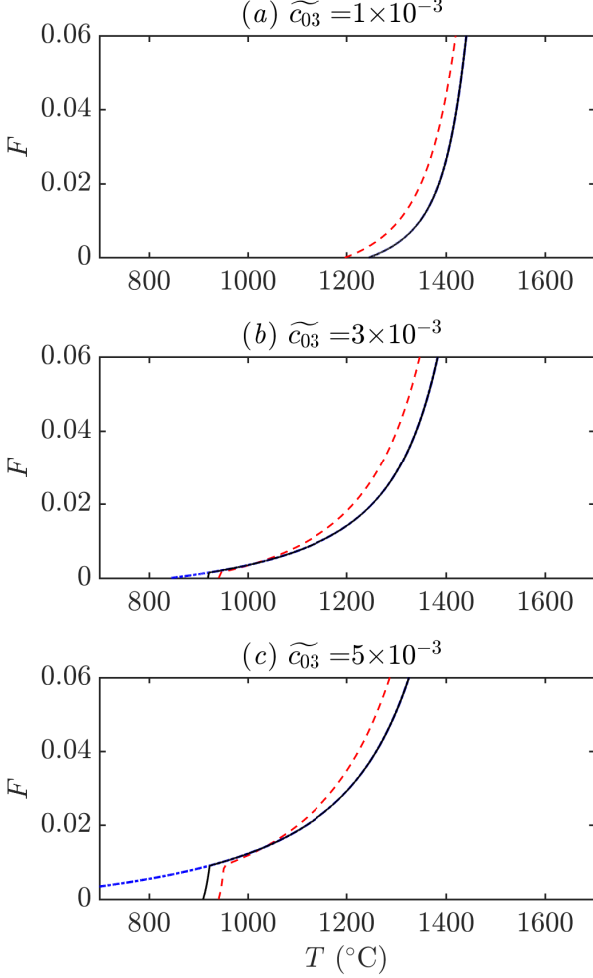


FIG. 6. The degree of melting $F(T)$ produced by our revised model (solid black) and the parameterization of Katz, Spiegelman, and Langmuir [18] (dashed red). Results computed at fixed pressure (3 GPa) at increasing water content: (a) $\widetilde{c}_{03} = 1 \times 10^{-3}$, (b) $\widetilde{c}_{03} = 3 \times 10^{-3}$, and (c) $\widetilde{c}_{03} = 5 \times 10^{-3}$. Note the kink in the curves around 950°C in (b, c), which is associated with water saturation. Without this saturation behaviour, the standard model predicts melting at several hundred degrees cooler temperatures (dashed blue curves).

II. FURTHER DETAILS OF ONE-DIMENSIONAL COLUMN MODEL

In the context of a one-dimensional melting model, conservation plays a strong role in constraining the model behaviour in steady state. We adopt an extended Boussinesq approximation in which we neglect density differences between the phases except for their role in driving fluid flow buoyantly. There are several equivalent ways to present the following equations; we approach the problem by considering conservation in the liquid phase and in the two-phase composite.

Mass conservation gives

$$\frac{d}{dz}(\phi w^l) = \frac{\Gamma}{\rho}, \quad (21)$$

$$\frac{d}{dz}\bar{w} = 0, \quad (22)$$

where $\bar{x} = x^s(1 - \phi) + x^l\phi$ denotes an average over the solid and liquid phases, with volume fractions $(1 - \phi)$ and ϕ respectively. The vertical velocity is w , volumetric melting rate is Γ and density is ρ . We first integrate equation (22) to obtain

$$\frac{\phi w^l}{\bar{W}_0} + \frac{(1 - \phi)w^s}{\bar{W}_0} = 1, \quad (23)$$

where \bar{W}_0 is the volume flux at the bottom of the melting column (which is not the motion of the solid phase alone, unlike in upwelling mantle columns used in the context of mid-ocean ridge magmatism). We follow the approach of Ribe [14] and define the quantity $F = \phi w^l / \bar{W}_0$. Thus the scaled liquid phase volume flux is F and the scaled solid phase volume flux is $(1 - F)$.

We can recover our previous definition of F in equation (8) by considering conservation of species mass. For each component $j = 1, 2, 3$,

$$\frac{d}{dz}(\phi w^l c_j^l) = \frac{\Gamma_j}{\rho}, \quad (24)$$

$$\frac{d}{dz}\bar{w}c_j = 0. \quad (25)$$

Note that, by summing equation (24) over j and comparing with equation (21), $\sum_j \Gamma_j = \Gamma$. We integrate equation (25) and use equation (23) to obtain

$$F c_j^l + (1 - F) c_j^s = \frac{\bar{W}_0 c_{0j}}{\bar{W}_0} \equiv \widetilde{c}_{0j}. \quad (26)$$

We then determine the degree of melting F , which is controlled by an energy equation and our phase diagram. One unusual feature of subduction zones is the non-monotonic temperature profile, which is largely controlled by the flow of the solid mantle. We use a steady energy balance for a one-dimensional column

$$\rho c_p \bar{W}_0 \frac{dT}{dz} = -L\Gamma + \frac{d}{dz} \left(\rho c_p \kappa \frac{dT}{dz} \right) + \rho c_p \Psi, \quad (27)$$

with a volumetric source term $\rho c_p \Psi$ that represents horizontal advection and diffusion of the background solid flow. In the absence of melting, the final pair of terms on the right-hand side establishes a non-monotonic temperature profile.

Next we observe that $\bar{W}_0 F' = \Gamma / \rho$ and F' is proportional to the isobaric productivity discussed previously, namely $F' = (T' + \rho g / \gamma) \partial F / \partial T$. We can better understand the system by non-dimensionalizing the energy equation. We scale lengths by H (the depth of the

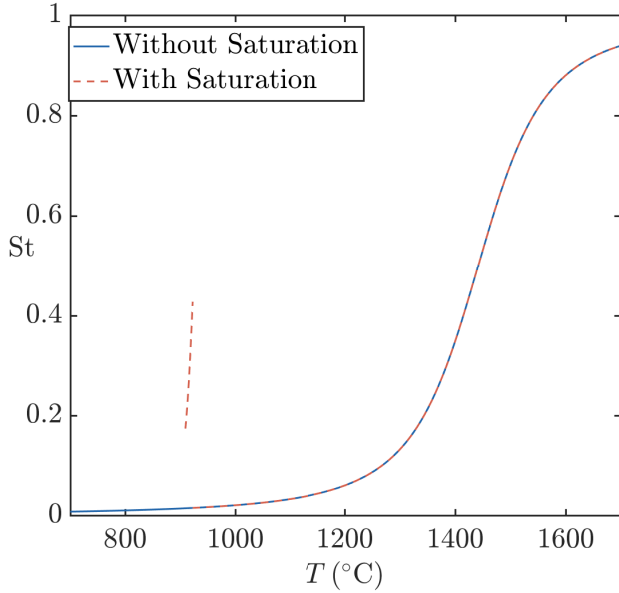


FIG. 7. Stefan number $St = (L/c_p)\partial F/\partial T$ at $P = 3$ GPa with and without modified phase diagram to account for water saturation, as discussed in section IC. Note that there is now a small segment of higher productivity between the solidus temperature and the temperature at which the melt becomes saturated.

melting column), and the source term by κ/H^2 . The dimensionless parameters involved are a Péclet number $Pe = HW_0/\kappa$, a Stefan number $St = (L/c_p)\partial F/\partial T$, a temperature change $\Delta T_H = \rho g H/\gamma$. Then the energy equation is

$$T'' = -\Psi + Pe [T'(1 + St) + St \Delta T_H], \quad (0 \leq z \leq 1). \quad (28)$$

A scaled version of the melting rate is

$$\tilde{\Gamma} \equiv \frac{H^2}{\kappa} \frac{\Gamma}{\rho} = Pe \frac{\partial F}{\partial T}(T' + \Delta T_H), \quad (29)$$

which has units of degrees Kelvin. Equation (28) is subject to boundary conditions on T at $z = 0$ and $z = 1$. In general, the Péclet number is fixed but the Stefan number depends on temperature and pressure (hence depth), as well as the compositional parameters of our melting model. We plot the Stefan number in Fig. 7.

Our experimental approach is as follows. Extract a vertical temperature profile T_{ref} from a single phase mantle flow thermal model of a subduction zone, as shown in Fig. 8. We then calculate $\Psi = -T''_{\text{ref}}(z)$. To investigate the effect of melting, we solve the dimensionless energy equation, focussing on the effect of varying the Péclet number and Stefan number (since the decompression term ΔT_H is well known).

We show a set of typical results in the main article. The main physical mechanism giving rise to this temperature signal is the fluid advection term in equation (28). Indeed, the latent heat of freezing/melting is

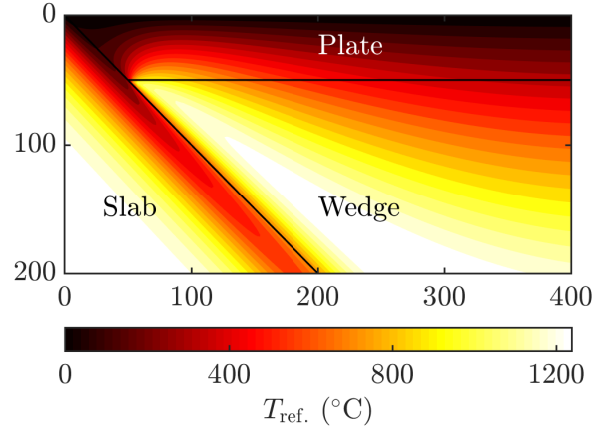


FIG. 8. Reference temperature field T_{ref} from the benchmark of van Keken *et al.* [19]. The dip angle, slab velocity and thickness of the overriding plate are prescribed. The solid velocity in the mantle wedge is calculated and coupled to the temperature through the temperature-weakening viscosity. We show only a subset of the full computational domain, which is 660 km wide and 600 km deep.

rather small and can be neglected. In Fig. 7 we see that $S < 0.1$ throughout the temperature range encountered ($T_{\text{ref}} \leq 1250^\circ\text{C}$), and in Fig. 9 we show that neglecting latent heat makes very little difference to the temperature profiles. One possible concern with this conclusion is that our standard phase diagram does not have an increased region of productivity associated with saturation of the melt phase, as discussed in section IC. This increased productivity corresponds to a significantly increased Stefan number for temperatures between the solidus temperature and the temperature at which the melt is saturated in water (Fig. 7). However, this increased productivity only occurs over a narrow temperature range and is not significant in changing the thermal profile predicted (Fig. 9).

III. FURTHER DETAILS OF TWO-DIMENSIONAL THERMAL MODEL

Sources of fluids in subduction zones are believed to be localized to particular depth ranges, associated with particular dehydration reactions in the subducting slab. Thus, in addition to the calculations presented in the main article, we also take three Gaussian fluid flow profiles associated with the major dehydration reactions of the slab, with a position, magnitude and width suggested by Wilson *et al.* [20]. We also consider the effect of all three sources combined.

As before, the principal result of our thermal calculations is that fluid flow can indeed substantially alter the thermal fluid in subduction zones, as shown in Fig. 10. Fluid flow associated with the peridotite source (a) is

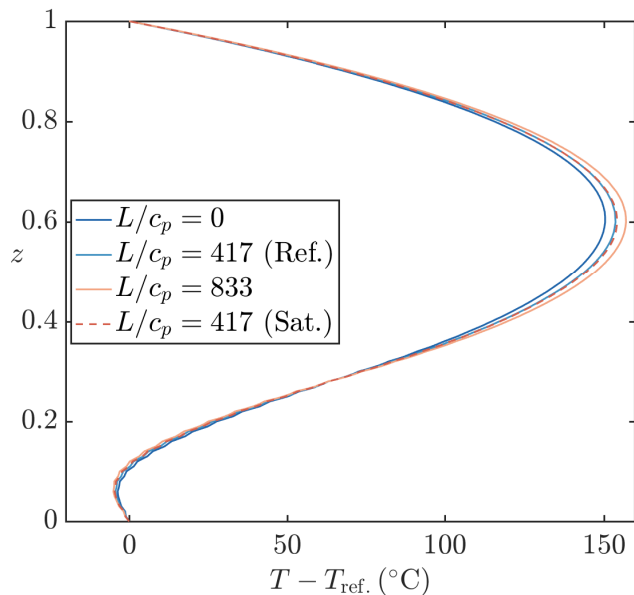


FIG. 9. Thermal structure is not significantly affected by latent heat release. We show temperature change with variations to the reference form of latent heat release in the energy equation (28) at fixed $Pe = 1$. Units are ($^{\circ}C$). Note that the Stefan number $St = (L/c_p)\partial F/\partial T$. Other parameters are as in Fig. 7.

the most thermally significant, raising temperatures by over 200 K. Fluid flow associated with the MORB source (*b*) raises temperatures near the trench by about 40 K; the gabbro source (*c*) is thermally insignificant. The peridotite source is strongest because it is associated with the largest fluid flow. The gabbro source is especially weak because it is rather narrower than the other sources, and so tends to diffuse laterally more strongly. The combined set of sources (*d*) is dominated by the peridotite source, although there are also slightly elevated temperatures in the fore-arc region associated with the MORB source.

We next consider the physical mechanisms that affect the thermal perturbation associated with magmatism. For all these calculations we return to the arc case with reference magma flux from the main article.

First, we show that the total amount of magma flux is more significant than the width of the source. In Fig. 11, we show that similar temperatures are found with the same total flux but very different widths. The width of the thermal response is controlled primarily by the balance between advective heat transport by the magma and thermal diffusion.

Second, we consider the effect of the coupling between the solid velocity and the temperature field through the temperature-weakening viscosity. We perform semi-decoupled calculations in which the solid velocity is fixed at the reference conditions associated with the reference temperature field (i.e., that without magmatism). In Fig. 12, we show that the fully coupled calculations are substantially warmer than the semi-decoupled calculations.

The mechanism behind this change is the increased circulation in the mantle wedge, leading to increased heat transport, shown in Fig. 13. The effect of coupling is more pronounced with smaller plate thickness, because there is a larger region of mantle flow where the viscosity is reduced, leading to faster circulation.

Third, we consider the effect of plate thickness. The thermal effect of magmatism decreases slightly with increasing plate thickness, as shown in Fig. 14. This is associated with cooler temperatures in the reference state, reducing the advection of heat by the magma. The decrease is also aided by the fact that the coupling to the solid velocity becomes a less significant positive feedback as plate thickness increases.

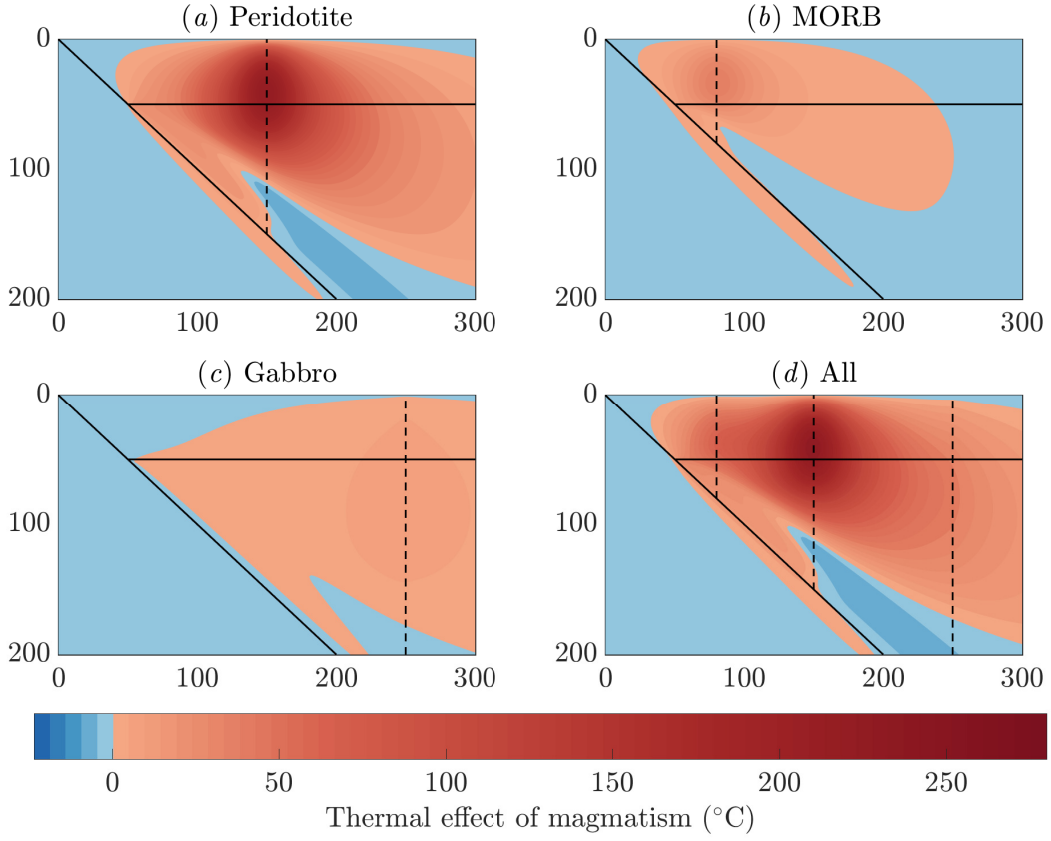


FIG. 10. The thermal impact of magmatism $T - T_{\text{ref}}$. (units $^{\circ}\text{C}$) associated with the dehydration of (a) peridotite, (b) MORB, and (c) gabbros. We also show (d) results when all three sources are combined. We indicate the individual sources as dashed lines at the centre of each Gaussian pulse. Horizontal and vertical scales are distance from the trench (units km).

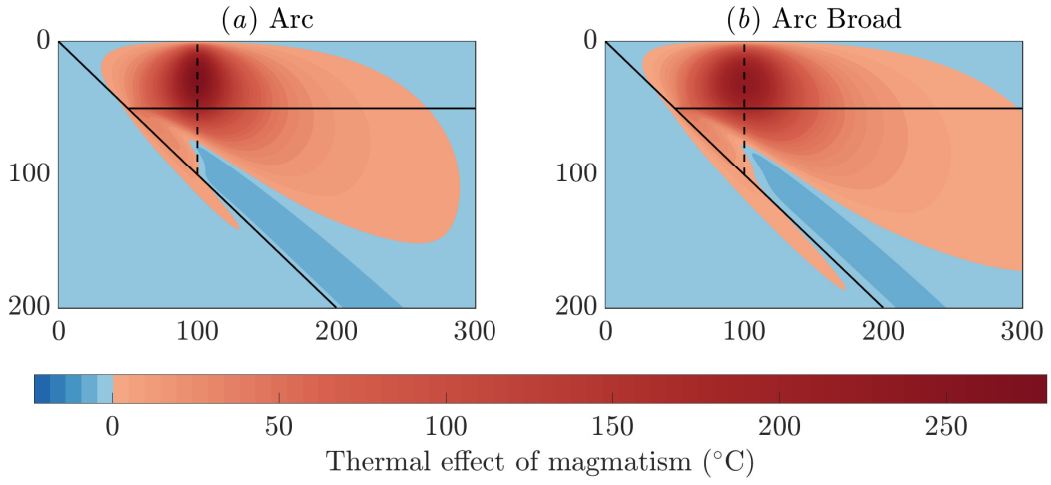


FIG. 11. Effect of width of the fluid source. (a) The standard reference arc case. (b) The arc case but with a source double the width but half the magnitude. The total flux is identical and the thermal perturbation is similar. The broad case has a slightly lower peak (by 40°C) and is slightly more diffuse. However, these differences are minor compared to those associated with varying the total magma flux as shown in the main article.

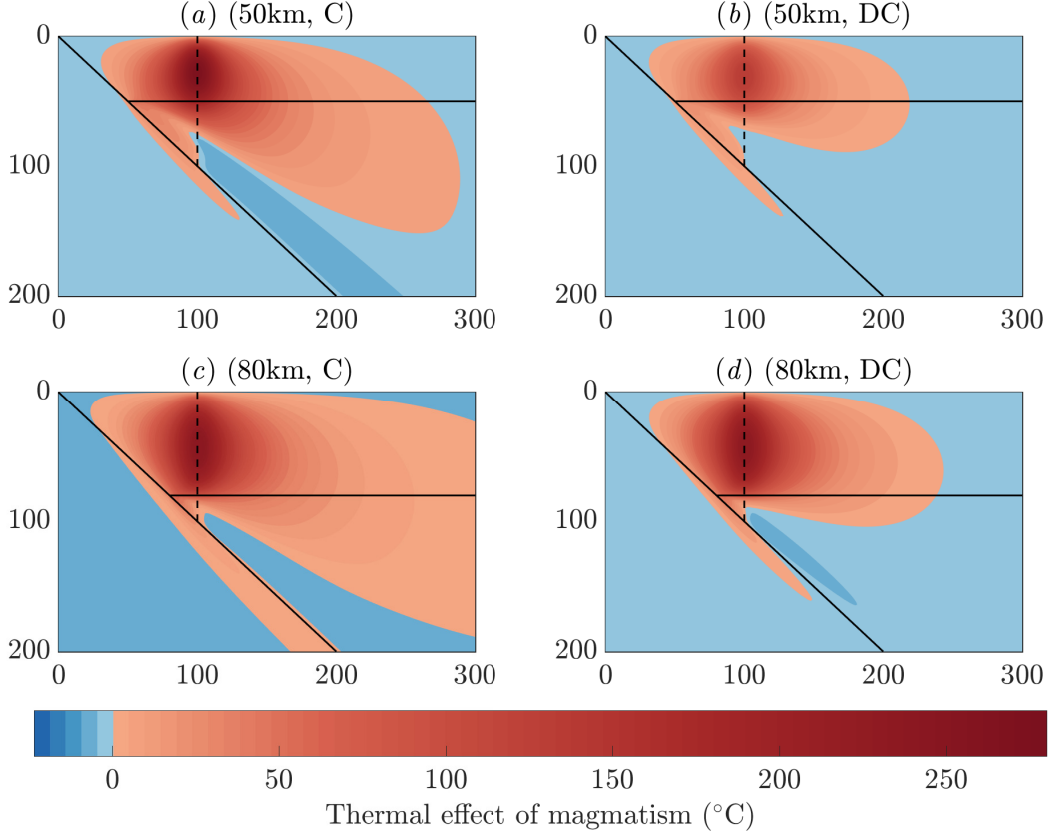


FIG. 12. Effect of changes in solid flow and plate thickness. We perform experiments in which we vary the thickness of the overriding plate (either 50 km or 80 km) and whether the solid flow is coupled to the perturbed temperature field (or just the reference field) denoted C (coupled) or DC (semi decoupled).

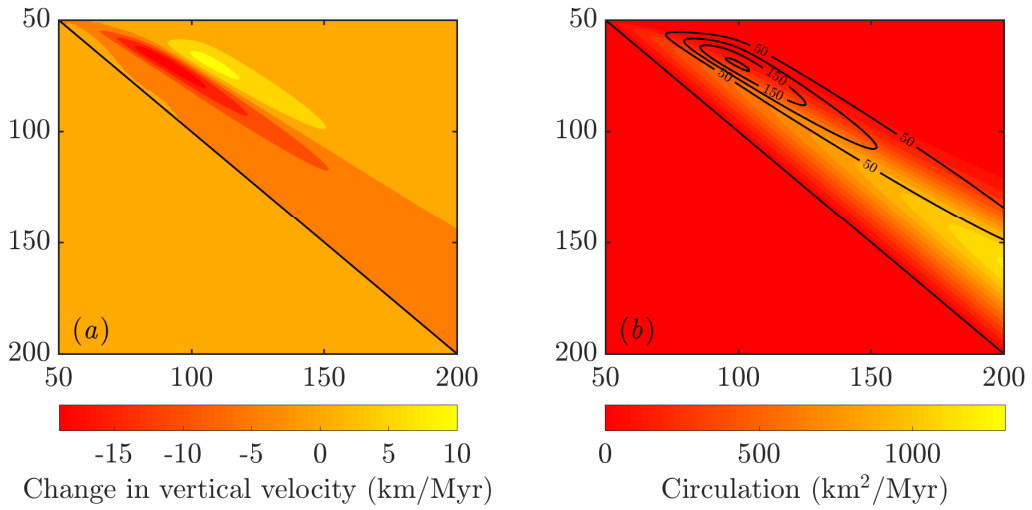


FIG. 13. Change in solid velocity associated with the thermal impact of magmatism. (a) changes in vertical velocity, which are moderately significant compared to the speed of the subducting slab which is 50 km/Myr. (b) circulation (streamfunction) is shown as the colour scale, with solid contours showing the change in the circulation due to the thermal impact of magmatism.

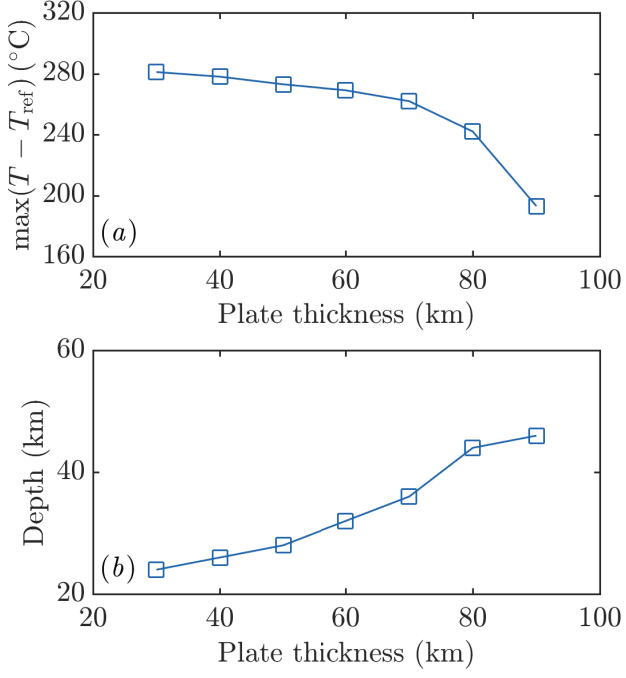


FIG. 14. Effect of the prescribed thickness of the overriding plate (reference arc case). (a) Maximum thermal perturbation due to magmatism, and (b) the depth at which the maximum occurs.